

K761223 WIDEX HEARING AID MODEL F8Dec 16, 1976
8 days to decisionK761223 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k761223/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Dec 8, 1976
Decision date	Dec 16, 1976
Days to decision	8 days
Third-party review	No

APPLICANT

Company	Widex Hearing Aid Co., Inc.
Location	Mchenry, IL, US
510(k) history	52 submissions · 52 cleared · 1976-2008

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k761223/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 25, 2026